

**DISPERSIONS OF INHERENTLY CONDUCTIVE POLYMER
IN NON-IONIC WATERBORNE POLYMERS**

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Claim of Priority

This application claims priority from U.S. Provisional Patent
Application Serial Number 60/520,026 bearing Attorney Docket Number
10 12003025 and filed on November 14, 2003.

Field of the Invention

This invention relates to dispersions of inherently conductive polymer in
waterborne polymers.

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Background of the Invention

Inherently conductive polymers have been found to be very useful in a
variety of electrically active materials, such as anti-fouling marine coatings;
anti-static fabrics, coatings and packaging; batteries; conductive inks;
20 conductive adhesives; EMI/RFI shielding articles, radar or microwave
absorption articles, and sensors.

Inherently conductive polymers can be the matrix of the coating. More
preferably because of cost, inherently conductive polymers are dispersed in an
inert binder to serve as the matrix or continuous phase of the coating.

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Summary of the Invention

What the art needs is a dispersion of inherently conductive polymers in a
binder that is environmentally friendly, relatively inexpensive, easy to apply,
and good performing.

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The present invention solves the problem in the art by providing a
mixture of inherently conductive polymer in a non-ionic waterborne polymer,

preferably a non-ionic waterborne polymer containing acrylic monomer or urethane monomer. More preferably, the non-ionic polymer is an aqueous "breathable" polyurethane binder.

5 Most particularly, the present invention uses a polyurethane binder that is an aliphatic polyether waterborne urethane polymer that has a high moisture vapor transmission rate (MVTR).

One aspect of the invention is a coatable mixture comprising inherently conductive polymer and a non-ionic waterborne polymer.

10 A feature of the invention is that the non-ionic polymer does not interfere with the conductive properties of the inherently conductive polymer.

An advantage of the present invention is the coatable mixture is stable under conventional storage and application conditions, is capable of renewing conductive polymeric properties of the inherently conductive polymer because the preferred non-ionic polyurethane has a high MVTR that causes rapid
15 absorption of moisture by the conductive polymer which is important for good conductivity.

Other advantages of the invention will become apparent when considering the embodiments of the invention.

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Embodiments of the Invention

Inherently Conductive Polymers

Inherently conductive polymers suitable for the invention include
25 polymers having repeating monomeric units of aniline, thiophene, pyrrole, phenyl mercaptan, and the like. Other examples include a conducting polymer selected from the group consisting of substituted and unsubstituted polyparaphenylenevinylenes, substituted and unsubstituted polyanilines, substituted and unsubstituted polyazines, substituted and unsubstituted
30 polythiophenes, substituted and unsubstituted polyparaphenylenes, substituted and unsubstituted poly-p-phenylene sulfides, substituted and unsubstituted

polyfuranes, substituted and unsubstituted polypyrroles, substituted and unsubstituted polyselenophene, substituted and unsubstituted polyacetylenes, mixtures thereof, and copolymers thereof. These conductive polymers are disclosed in a variety of patents, including U.S. Pat. Nos. 5,069,820 (Jen et al.); 5,160,457 (Elsenbaumer); 5,185,100 (Han et al.); 5,281,363 (Shacklette et al.); 5,378,403 (Shacklette); 5,422,423 (Shacklette et al.); 5,456,862 (Kwan-Yue et al.); 5,567,355 (Wessling et al.); 5,700,398 (Angelopoulos et al.) and 5,911,918 (Shacklette et al.). As described in these patents the inherently conductive polymer is often doped with an acid such as hydrochloric acid or p-toluene sulfonic acid.

Particularly preferred is a substituted polyaniline such as disclosed in U.S. Pat. No. 5,968,417 (Visawanathan) and more particularly that marketed by PolyOne Corporation as Teslart™ inherently conductive polymers. This substituted polyaniline is lignosulfonic acid-grafted polyaniline.

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Non-Ionic Waterborne Polymers

Waterborne polymers are useful in the present invention because of the desire in the industry to avoid organic solvents. Non-ionic waterborne polymers provide a stable environment for the operation of the inherently conductive polymer. It has been found that ionic waterborne polymers, either cationic or anionic, adversely affect the conductivity of the inherently conductive polymer.

Commercially available non-ionic waterborne polymers are known to contain polymers with either acrylic monomers or urethane monomers. This type of polymer is emerging in industry as a useful non-ionic carrier for a variety of specialized additives for the coatings industry. As such new non-ionic waterborne polymers become available, one skilled in the art without undue experimentation will be able to determine the suitability of such new products for use in the present invention.

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Polyurethane

Non-ionic polyurethanes are preferred for the present invention, especially those which are called "breathable" polyurethanes because they exhibit high MVTR properties.

Breathable polyurethanes comprise (a) poly(alkylene oxide) side-chain units in an amount comprising about 12 wt. % to about 80 wt. % of the polyurethane, wherein (i) alkylene oxide groups in said poly(alkylene oxide) side-chain units have from 2 to 10 carbon atoms and are unsubstituted, substituted, or both unsubstituted and substituted, (ii) at least about 50 wt. % of said alkylene oxide groups are ethylene oxide, and (iii) the amount of side-chain units is (i) at least about 30 wt. % when the molecular weight of side-chain units is less than about 600 grams/mole, (ii) at least about 15 wt. % when the molecular weight of side-chain units is from about 600 to about 1,000 grams/mole, and (iii) at least about 12 wt. % when the molecular weight of side-chain units is more than about 1,000 grams/mole, and (b) poly(ethylene oxide) main-chain units in an amount comprising less than about 25 wt. % of the polyurethane.

Such breathable polyurethanes are disclosed in detail in United States Patent Publication 20030195293 (Lubnin et al.).

Non-limiting commercial examples of polyurethanes suitable for the invention include Permax™ 200 and 220 urethane emulsions available from Noveon, Inc. of Brecksville, Ohio. Additional information can be found at www.noveoncoatings.com.

Mixing of Inherently Conductive Polymers and Waterborne Polymers

Conventional mixing equipment is used to thoroughly mix the inherently conductive polymer into the non-ionic polymer emulsion.

The amount of inherently conductive polymer added to the polymer emulsion can range from about 5 to about 50 weight percent of total solids, and preferably from about 10 to about 25 weight percent of total solids of the mixture.

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Optional Ingredients

A variety of ingredients commonly used in the coatings industry can also be included in the mixture of the present invention. Non-limiting examples of such optional additives include slip agents, antiblocking agents, antioxidants, ultraviolet light stabilizers, quenchers, plasticizers, lubricants, antistatic agents, fire retardants, and fillers such as glass fibers, talc, chalk, or clay. Of these fillers, the properties of nanoclay can add stiffness, toughness, and charring properties for flame retardancy. Such optional additives can be included in the mixture of the present invention in an amount from about 0 to about 80, and preferably from about 0.1 to about 50 weight percent. Most preferably, the amount is about 1 to about 30 weight percent of the total solids of the mixture.

Any conventional colorant useful in coatings and paints is also acceptable for use in the present invention. Conventional colorants can be employed, including inorganic pigments such as titanium dioxide, iron oxide, chromium oxide, lead chromate, carbon black, silica, talc, china clay, metallic oxides, silicates, chromates, etc., and organic pigments, such as phthalocyanine blue, phthalocyanine green, carbazole violet, anthrapyrimidine yellow, flavanthrone yellow, isoindoline yellow, indanthrone blue, quinacridone violet, perylene reds, diazo red and others. The amount of colorant can range from none at all to about 30, and preferably from about 1.5 to about 10 weight percent of total solids of the mixture.

An especially desired optional ingredient is a non-ionic thickener or anti-settling agent to promote better retained dispersion of the inherently conductive polymer particles in the non-ionic waterborne polymer after initial mixing and when coating the mixture in larger film thicknesses. Commercially available thickeners or rheology modifiers are sold by Elementis Specialties of Hightstown, NJ under the Rheolate brand, some of which are believed to be non-ionic associative urethane thickeners. More information can be found at www.elementis-specialties.com. The amount of thickener can range

from about none at all to about 5, and preferably from about 1 to about 3 weight percent of total solids of the mixture.

Usefulness of the Invention

5 All of the advantages and usefulness of a breathable urethane polymer as disclosed in United States Patent Publication 20030195293 (Lubnin et al.) are also present in the mixture of the present invention. But the inherently conductive polymer ingredient adds to those properties by making the mixture electrically active and capable of serving as an electron transfer agent.

10 Surface resistivities, using the Four Point Probe test (ASTMD-257-99) can range from about 1.00×10^3 to about 1.00×10^{12} Ohms/square.

The coatable mixture can be prepared with approximately 30% solids and a Brookfield viscosity of about 125 Centapoise. This permits ease of application to a substrate via spray, brush, roll, knife, or other means of
15 application.

Thus, mixtures of the present invention can be very useful in a variety of electrically active materials, such as anti-fouling marine coatings; anti-static fabrics, coatings and packaging; batteries; conductive inks; conductive adhesives; EMI/RFI shielding articles, radar or microwave absorption articles,
20 and sensors.

While not being limited to a particular theory, it is believed that the non-ionic nature of waterborne polymers used in the present invention inhibits coagulation of the mixture that might otherwise be caused by protons present in the mixture due to the acidic nature of the inherently conductive polymer.

25 Moreover, it is believed that the high MVTR of the commercial Permax™ polyurethane material promotes rapid re-absorption of water from the environment by the inherently conductive polymer, which is important to good conductivity values for the inherently conductive polymer, after the mixture of the present invention is coated on a surface and dried.

30 Further embodiments are described in the following examples.

Examples

Table 1 shows the commercial ingredients used in Examples 1-5 and Comparison Example A.

Table 1 -- Source of Ingredients					
Ingredient Name	Purpose	Brand Name	Generic Name	Source	Source Location
Bayhydrol 110	Polymeric Binder	Bayhydrol	Ionic Waterborne Polyurethane Dispersion	Bayer	Pittsburgh, PA
Permax 220	Polymeric Binder	Permax	Non-Ionic Waterborne Polyurethane Dispersion	Noveon	Brecksville, OH
Teslart™ Wet Cake	Conductive Filler	Teslart™	Polyaniline chains grafted to ligno-sulfonic acid	PolyOne Corp.	Avon Lake, OH
Rheolate 300	Thickener Anti Settling Agent	Rheolate	Non-Ionic Thickener	Elementis Specialties	Hightstown, NJ

Table 2 shows the Recipes, Method of Preparation, and resulting Properties of Examples 1-5 of mixtures of the present invention, in comparison with Comparison Example A, which uses a polyurethane without high MVTR.

Table 2						
Recipes, Preparation, and Properties						
Ingredient	Example 1	2	3	4	5	A
Recipes						
Bayhydrol 110	0%	0%	0%	0%	0%	89.0%
Permamax 220	89.0%	87.2%	78.8%	71.8%	66.0%	0%
Teslart™ Wet Cake (28.5% Solids)	11.0%	10.7%	19.3%	26.5%	32.4%	11.0%
Rheolate 300	0%	2.1%	1.9%	1.7%	1.6%	0%
Preparation						
Mixing Equipment	Marine Prop	Marine Prop	Marine Prop	Marine Prop	Marine Prop	Marine Prop
Mixing Temp.	Room	Room	Room	Room	Room	Room
Mixing Speed	200-500 RPM	200-500 RPM	200-500 RPM	200-500 RPM	200-500 RPM	200-500 RPM
Order of Addition of Ingredients	Permamax & Teslart, mixed well	Permamax & Teslart, mixed well, then Rheolate	Permamax & Teslart, mixed well, then Rheolate	Permamax & Teslart, mixed well, then Rheolate	Permamax & Teslart, mixed well, then Rheolate	Bayhydrol & Teslart, mixed well
Form of Product After Mixing	Dispersion	Dispersion	Dispersion	Dispersion	Dispersion	Dispersion
Properties						
Time to Coagulation at Room Temp. (Days)	>90	>90	>90	>90	>90	<1
Surface Resistivity (ASTM D257-99) Ohms/sq (2 mils thick (0.05 mm) dry, 24hr @ 50% RH)	Not Measured – Too Much Settling	9 * 10E10	3 * 10E5	3 * 10E5	2 * 10E5	Not Measured – Dispersion Coagulated

Table 2 shows that all of Examples 1-5 using a non-ionic polyurethane do not coagulate within a day, as compared with use of an ionic polyurethane of Comparison A Example, does coagulate. Examples 3-5 perform better
5 concerning surface resistivity than Example 2, indicating that a thickener or anti-settling agent is preferable in the mixture of the present invention. Also, surface resistivity decreases as loading of inherently conductive polymer increases, as shown by Example 5 having a lower surface resistivity than Example 3.

10 The invention is not limited to the above embodiments. The claims follow.